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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
CA SUBSCRIBER PRICE	ENTRY -1.60	SESSION -1.60
=> file registry COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	11.35	259.57
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-1.60	-1.60

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STRUCTURE FILE UPDATES: 21 JAN 2008 HIGHEST RN 1000370-19-3 DICTIONARY FILE UPDATES: 21 JAN 2008 HIGHEST RN 1000370-19-3

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TSCA INFORMATION NOW CURRENT THROUGH June 29, 2007

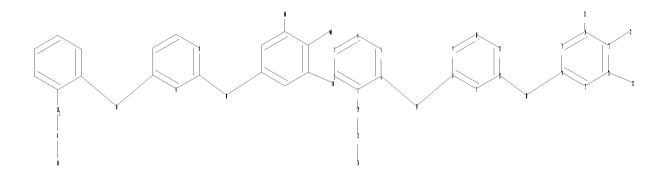
Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=>

Uploading C:\Program Files\STNEXP\Queries\10571733.str



chain nodes : 19 20 21 22 23 24 25 26 ring nodes : $1 \quad 2 \quad 3 \quad 4 \quad 5 \quad 6 \quad 7 \quad 8 \quad 9 \quad 10 \quad 11 \quad 12 \quad 13 \quad 14 \quad 15 \quad 16 \quad 17 \quad 18$ chain bonds : 1-24 6-19 8-19 12-20 14-20 16-21 17-22 18-23 24-25 25-26ring bonds : $1-2 \quad 1-6 \quad 2-3 \quad 3-4 \quad 4-5 \quad 5-6 \quad 7-8 \quad 7-12 \quad 8-9 \quad 9-10 \quad 10-11 \quad 11-12 \quad 13-14 \quad 13-18$ 14-15 15-16 16-17 17-18 exact/norm bonds : 6-19 8-19 12-20 14-20 24-25 25-26 exact bonds : 1-24 16-21 17-22 18-23 normalized bonds : $1-2 \quad 1-6 \quad 2-3 \quad 3-4 \quad 4-5 \quad 5-6 \quad 7-8 \quad 7-12 \quad 8-9 \quad 9-10 \quad 10-11 \quad 11-12 \quad 13-14 \quad 13-18$ 14-15 15-16 16-17 17-18

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS

L5 STRUCTURE UPLOADED

=> 15 sss ful

L5 IS NOT A RECOGNIZED COMMAND

The previous command name entered was not recognized by the system. For a list of commands available to you in the current file, enter "HELP COMMANDS" at an arrow prompt (=>).

=> s 15 sss ful FULL SEARCH INITIATED 12:53:21 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 66 TO ITERATE

100.0% PROCESSED 66 ITERATIONS 5 ANSWERS SEARCH TIME: 00.00.01

L6 5 SEA SSS FUL L5

=> d 16 1-6

L6 ANSWER 1 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN

RN 848468-37-1 REGISTRY

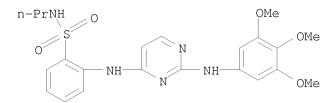
ED Entered STN: 14 Apr 2005

CN Benzenesulfonamide, N-propyl-2-[[2-[(3,4,5-trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)

MF C22 H27 N5 O5 S

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L6 ANSWER 2 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN

RN 848468-35-9 REGISTRY

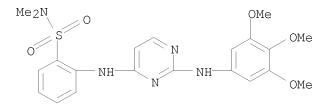
ED Entered STN: 14 Apr 2005

CN Benzenesulfonamide, N,N-dimethyl-2-[[2-[(3,4,5-trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)

MF C21 H25 N5 O5 S

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L6 ANSWER 3 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN

RN 848468-34-8 REGISTRY

ED Entered STN: 14 Apr 2005

CN Benzenesulfonamide, N-(2-hydroxyethyl)-2-[[2-[(3,4,5-trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)

MF C21 H25 N5 O6 S

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L6 ANSWER 4 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN
- RN 604802-45-1 REGISTRY
- ED Entered STN: 15 Oct 2003
- CN Benzenesulfonamide, N,2-dimethyl-6-[[2-[(3,4,5-trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)
- MF C21 H25 N5 O5 S
- SR CA
- LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
- 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L6 ANSWER 5 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN
- RN 604802-41-7 REGISTRY
- ED Entered STN: 15 Oct 2003
- CN Benzenesulfonamide, N-methyl-2-[[5-(trifluoromethyl)-2-[(3,4,5-trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)
- MF C21 H22 F3 N5 O5 S
- SR CA
- LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file caplus
COST IN U.S. DOLLARS

SINCE FILE TOTAL
ENTRY SESSION
188.82 448.39

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL
ENTRY SESSION

0.00

-1.60

FILE 'CAPLUS' ENTERED AT 12:53:50 ON 22 JAN 2008
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=> s 16

L7 2 L6

CA SUBSCRIBER PRICE

=> d 17 ibib abs 1-2

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2005:260035 CAPLUS

DOCUMENT NUMBER: 142:336377

TITLE: Preparation of 2,4-di(phenylamino)pyrimidines useful

in the treatment of proliferative disorders

INVENTOR(S): Imbach, Patricia; Roesel, Johannes

PATENT ASSIGNEE(S): Novartis AG, Switz.; Novartis Pharma GmbH

SOURCE: PCT Int. Appl., 39 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.					KIND DATE			,	APPI		DATE							
WO	2005026130			A1 20050324				 WO 2	2004-	20040917								
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,	
											EC,							
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	ΝI,	
		NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
		ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW	
	RW:	BW,	GH,	GM,	ΚE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	
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											LU,							
		SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	ΝE,	
			TD,															
	AU 2004272288																	
CA	2538	38413			A1 20050324				CA 2	2004-		2	0040	917				
EP		1663992			A1 20060607													
	R:										ΙΤ,			ΝL,	SE,	MC,	PT,	
											HU,							
	CN 1852900				А		2006	1025		CN 2	2004-	8002	6942		2	0040		
BR	BR 2004014544						2006	1107		BR 2	2004-		20040917					
JP	JP 2007505858				T 20070315				JP 2006-526595									
US	US 2007105839				A1 20070510			US 2006-571733										
									IN 2006-CN922									
					A 20060531			MX 2006-PA3054										
RIORIT	Y APP	LN.	INFO	.:							2003-					0030		
											2004-				W 2	0040	917	
THER S	OURCE	(S):			CAS:	REAC	T 14	2:33	6377	; MA	RPAT	142	:336.	377				

GΙ

AB The title compds. I [X = CR0, N; R0, R1-R4 = H, OH, alkyl, etc.; or R3 and R4 form together with the nitrogen and carbon atoms to which they are attached a 5-10 membered heterocyclic ring and comprising addnl. 1-3 heteroatoms selected from N, O and S; or R1-R3 = halo, haloalkyl, alkoxy, etc.; or R1 and R2 form aryl or 5-10 membered heteroaryl; R5, R6 = H, halo, CN, alkyl, etc.; R7-R9 = H, OH, alkyl, etc.], useful for preventing or treating proliferative disorders such as a tumor disease, by inhibiting ALK activity, were prepared E.g., a 2-step synthesis of 2-[2-(1H-indazol-6-ylamino)-pyrimidin-4-ylamino]benzenesulfonamide, starting from 2-aminobenzenesulfonamide and 2,4-dichloropyrimidine, was given. The compds. I were tested for inhibition of ALK tyrosine kinase in various cellular assays (data were given for representative compds. I).

Ι

REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN L7

ACCESSION NUMBER: 2003:757684 CAPLUS

DOCUMENT NUMBER: 139:292258

TITLE: Pyrimidine derivatives

INVENTOR(S): Baenteli, Rolf; Zenke, Gerhard; Cooke, Nigel Graham; Duthaler, Rudolf; Thoma, Gebhard; Von Matt, Anette; Honda, Toshiyuki; Matsuura, Naoko; Nonomura, Kazuhiko;

Ohmori, Osamu; Umemura, Ichiro; Hinterding, Klaus;

APPLICATION NO.

DATE

Papageorgiou, Christos

DATE

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis Pharma G.m.b.H.

PCT Int. Appl., 45 pp. SOURCE:

CODEN: PIXXD2

KIND

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.

17111111 110.															DITTE				
WO	2003	0784	04		A1	_	20030925			WO 2003-EP2710					20030314				
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		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC	EE,	ES,	FΙ,	GB,	GD,	GE,	GH,		
		HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG	KP,	KR,	KΖ,	LC,	LK,	LT,	LU,		
		LV,	MA,	MD,	MK,	MN,	MX,	NΙ,	NO,	ΝZ	, OM,	PH,	PL,	PT,	RO,	RU,	SC,		
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CA	2479	133		A1 20030925					CA	2003-	2479	20030314							
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EP								EP 2003-744366											
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											, TR,								
								BR 2003-8461											
JP	2005	5275:	29		Т				JP 2003-576410										
CN	1697	830			А			CN 2003-806101											
	5351								NZ 2003-535109										
		2004CN02025																	
		2004PA09058					2005				2004-								
		2004004374									2004-								
	S 2006100227									2005-									
	2004				А		2006	0531			2004-				_	20060			
IORIT:	Y APP	LN.	INFO	.:							2002-								
										WO	2003-	EP27	10	1	W 2	20030.	314		
THER SOURCE(S):					MARI	MARPAT 139 29225													

Ι

OTHER SOURCE(S): MARPAT 139:292258

GΙ

The pyrimidine derivs. (I) are claimed, wherein X = -CR or -N, R, R1, R2, AΒ R3, R4 independently is H, OH, C1-8alkyl, C2-8alkenyl, C3-8cycloalkyl, C3-8cycloalkyl-C1-8alkyl, hydroxyC1-8alkyl, C1-8alkoxyC1-8alkyl, hydroxyC1-8alkoxyC1-8alkyl, arylC1-8alkyl which optionally may be substituted on the ring by OH, C1-8alkoxy, carboxy, C1-8alkoxycarbonyl or ${
m R3}$ and ${
m R4}$ form together with N and C atoms to which they are attached to a 5-10 membered heterocyclic ring containing 1, 2 or 3 heteroatoms of N, O or S; R1 and R2 form together with C atoms to which they are attached aryl of 5-10 membered heteroaryl moiety containing 1-2 heteroatoms of N, O, S; R and R6 independently is H, halo, CN, C1-8alkyl, haloC1-8alkyl, C2-8alkenyl, C2-8alkynyl, C3-8cycloalkyl, C3-8cycloalkylC1-8alkyl, C5-10arylC1-8alkyl,; R7, R8 and R9 is independently H, OH, C1-8alkyl, C2-8alkenyl, haloC1-8alkyl, C1-8alkoxy, C3-8cycloalkyl, C3-8cycloalkylC1-8, arylC1-8alkyl. disorders where ZAP-70 and/or Syk inhibition plays a role or caused by a malfunction of signal cascades connected with FAK. I are useful in disorders where ZAP-70 and/or Syk inhibition plays a role or caused by a malfunction of signal cascades connected with FAK. Pharmaceutical compns. containing I are claimed. REFERENCE COUNT: THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS 9 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT => d his (FILE 'HOME' ENTERED AT 11:53:56 ON 22 JAN 2008)

FILE 'REGISTRY' ENTERED AT 11:54:20 ON 22 JAN 2008

STRUCTURE UPLOADED

L2 0 S L1 FAM FUL L3 5 S L1 SSS FUL

FILE 'CAPLUS' ENTERED AT 11:55:10 ON 22 JAN 2008

L4 2 S L3

FILE 'REGISTRY' ENTERED AT 12:52:56 ON 22 JAN 2008

STRUCTURE UPLOADED

L6 5 S L5 SSS FUL

FILE 'CAPLUS' ENTERED AT 12:53:50 ON 22 JAN 2008

L7 2 S L6

=> logoff

L1

L5

ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:y

COST IN U.S. DOLLARS
SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST
8.22
456.61

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

CA SUBSCRIBER PRICE ENTRY SESSION -1.60 -3.20

STN INTERNATIONAL LOGOFF AT 12:56:45 ON 22 JAN 2008